DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
1401 Rockville Pike
Rockville, MD 20852-1448

February 8, 2000

Ms. Roberta Grigg
MEDITECH Medical Information Technology, Inc.
MEDITECH Circle
Westwood, MA 02090

Re: BK980046
Product: Magic Client Server, Version 5.1
Date Received: 30-DEC-98
Classification: Unclassified

Dear Ms. Grigg:

We have reviewed your 510(k) notification of intent to market the device referenced above and have determined that the device is substantially equivalent to devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments. You may, therefore, market the device, subject to the general controls provisions of the Federal Food, Drug, and Cosmetic Act that include requirements for registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

Please be advised that as discussed with you by telephone on January 27, 2000, the following changes need to be made to your blood component labels before you begin marketing the software.

1. On all labels, the statement "Rx Only" should be in bold type.

2. On all labels, the statement "Properly Identify Intended Recipient" should be on a single line.

3. On the label for Red Blood Cells, Leukocytes Reduced, the product code should be "04360" instead of "04460".

If your device has been classified into either class II (Special Controls) or class III (Premarket Approval), (see above), it may be subject to the above and additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. In addition, the Food and Drug Administration

(FDA) may publish further announcements concerning your device in the Federal Register. Please note that this response to your Premarket notification submission does not affect any obligation you might have under the Radiation Control for Health and Safety Act of 1968, or other Federal Laws or regulations.

This letter will allow you to begin marketing your device as described. An FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and permits your device to proceed to the market, but it does not mean that FDA approves your device. Therefore, you may not promote or in any way represent your device or its labeling as being approved by FDA. If you desire specific advice on promotional labeling and advertisement for your device, please contact our Advertising and Promotional Labeling Staff (HFM-602) at (301) 827-3028.

Sincerely,

Jay S. Epstein, M.D.
Director
Office of Blood Research and Review
Center for Biologics Evaluation and Research